

Al-free di-trioctahedral substitution in chlorite and a ferri-sudoite end-member

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(Received 01 November 2015; revised 30 April 2016; Guest editor: Atsuyuki Inoue)

ABSTRACT: A compilation of Fe³⁺-bearing chlorite analyses is used: (1) to investigate the Al-free di-trioctahedral (AFDT) substitution $2\text{Fe}^{3+} + \square = 3(\text{Mg}, \text{Fe}^{2+})$ in chlorite; and (2) to estimate the composition of a ferri-sudoite end-member $(\text{Si}_3\text{Al})[(\text{Fe}^{2+}, \text{Mg})_2\text{Fe}_2^{3+}\square\text{Al}]\text{O}_{10}(\text{OH})_8$ within the chlorite solid-solution domain. According to our observations, up to two Fe³⁺ cations might be allocated in the M2-M3 chlorite sites by the substitution AFDT, which does not involve Al. These unexpected observations were made possible by the development of μXANES techniques allowing *in situ* measurements of $X\text{Fe}^{3+}$ ($\text{Fe}^{3+}/(\text{Fe}^{2+} + \text{Fe}^{3+})$) in heterogeneous chlorite. Although further studies are required to confirm the crystallographic position of Fe³⁺ and refine its ionic/magnetic behaviour in chlorite, this development creates opportunities for developing new geothermometers.

KEYWORDS: chlorite, μXANES , geothermometry, sudoite.

Chlorite is a ubiquitous phyllosilicate composed of tetrahedral and octahedral layers according to the 14.2 Å 2:1:1 arrangement (*e.g.* Meunier, 2005). The tetrahedral sheet hosts tri- or tetravalent cations (Si^{4+} , Al^{3+} or rarely Ti^{4+} , Fe^{3+}), while the octahedral one is composed of di- or trivalent cations (usually Fe^{2+} , Mg^{2+} , Mn^{2+} , Al^{3+} and Fe^{3+}). The general formula of chlorite is: $R_{6-x-3y}^{2+}R_{x+2y}^{3+}\square_y(\text{Si}_{4-x}\text{R}_x^{3+})\text{O}_{10}(\text{OH})_8$, in which R^{2+} represents divalent cations, R^{3+} trivalent cations and \square octahedral vacancies (*e.g.* Wiewióra & Weiss, 1990). According to Bailey (1988), the chlorite structure is formed of two types of tetrahedral crystallographic sites (denoted T1 and T2) and two octahedral sites (denoted M1 and M2). Two other octahedral sites (M3 and M4) are located in the brucite

interlayer space. Al^{VI} or trivalent Fe^{3+} cations are located preferentially in the M4 site, while the divalent cations occupy the other sites (Bailey, 1988). There are three categories of chlorite in diagenetic and low- to high-grade metamorphic rocks: tri-trioctahedral (sum of octahedral cations close to 6 a.p.f.u), di-trioctahedral (5 a.p.f.u) and di-dioctahedral (4 a.p.f.u). The main chemical substitutions occurring in chlorite are: Tschermak (TK) $\text{Si}_{(\text{IV})}^{4+} + (\text{Fe}^{2+}, \text{Mg}^{2+})_{(\text{VI})} = \text{Al}_{(\text{IV})}^{3+} + \text{Al}_{(\text{VI})}^{3+}$; ferromagnesian (FM) $\text{Mg}_{(\text{VI})}^{2+} = \text{Fe}_{(\text{VI})}^{2+}$ and di-trioctahedral (DT) $2\text{Al}_{(\text{VI})}^{3+} + \square_{(\text{VI})} = 3(\text{Mg}^{2+}, \text{Fe}^{2+})_{(\text{VI})}$. Two filling models are used in the literature for the distribution of cations on the crystallographic sites. The first model referred to as ‘disorderly’, considers that there is no preferred configuration (*e.g.* Inoue *et al.*, 2009). The second model, referred to as ‘ordered’ is constrained by specific rules for filling cations in the crystallographic sites such as preferred configuration and Fe-Mg equipartition (*e.g.* Vidal *et al.*, 2001, 2005, 2006; Lanari *et al.*, 2014a). These authors assume, following Holland *et al.* (1998), Al_{IV} in T2, vacancies in M1, Al_{VI}

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[†]This work was originally presented during the session ‘The many faces of chlorite’, part of the Euroclay 2015 conference held in July 2015 in Edinburgh, UK.

DOI: 10.1180/claymin.2016.051.4.09

TABLE 1. Structural formulae for classic chlorite end-members and the new ferri-sudoite end-member.

	T1(2)	T2(2)	M1(1)	M2-M3(4)	M4(1)
Amesite-Mg/corundophilite	Si,Si	Al,Al	Al	Mg,Mg,Mg,Mg	Al
Amesite-Fe	Si,Si	Al,Al	Al	Fe,Fe,Fe,Fe	Al
Clinochlore	Si,Si	Si,Al	Mg	Mg,Mg,Mg,Mg	Al
Chamosite/daphnite	Si,Si	Si,Al	Fe	Fe,Fe,Fe,Fe	Al
Sudoite-Mg	Si,Si	Si,Al	□	Mg,Mg,Al,Al	Al
Sudoite-Fe	Si,Si	Si,Al	□	Fe,Fe,Al,Al	Al
Al-free chlorite	Si,Si	Si,Si	Mg	Mg,Mg,Mg,Mg	Mg
Pyrophyllite-gibbsite	Si,Si	Si,Si	□	Al,Al,Al,□	Al
Ferri-sudoite	Si,Si	Si,Al	□	(Fe,Mg) ₂ ,Fe ³⁺ ,Fe ³⁺	Al

The numbers in parentheses refer to the quantity of atoms (per structural formula) distributed in tetrahedral (T) or octahedral (M) sites.

in M4 then M1 and then, eventually, in M2-M3, Fe and Mg in M2-M3 then in M1. Following the equipartition assumption, $\chi\text{Mg} \text{ (Mg}^{2+}/(\text{Mg}^{2+} + \text{Fe}^{2+}))$ is considered to be equal between the M1 and M2-M3 sites. Several chlorite end-members are classically defined (Table 1): Al-free chlorite (Si₄)[Mg₆]O₁₀(OH)₈, amesite (Si₂Al₂)[(Fe²⁺,Mg)₄Al₂]O₁₀(OH)₈, clinochlore (Si₃Al)[Mg₅Al]O₁₀(OH)₈, corundophilite (Si₂Al₂)[Mg₄Al₂]O₁₀(OH)₈, daphnite (Si₃Al)[Fe₅²⁺Al]O₁₀(OH)₈, pyrophyllite-gibbsite (Si₄)[Al₄□₂]O₁₀(OH)₈ and sudoite (Si₃Al)[(Fe²⁺,Mg)₂□Al₃]O₁₀(OH)₈. These end-members do not include any ferric iron component.

For 15 years, many authors have emphasized the importance of Fe³⁺ in chlorite, particularly in terms of its use as a geothermometer (Laird, 1988; Vidal *et al.*, 2006; Inoue *et al.*, 2009; Bourdelle *et al.*, 2013; Lanari *et al.*, 2014a). An additional substitution: Al-Fe³⁺ that is supposed to occur on the M4 site was defined but without defining a ferric chlorite end-member (Vidal *et al.*, 2006). In this model, the incorporation of Fe³⁺ changes the chlorite composition and the activity of the other end-members. This choice of the M4 site is based on the results of Smyth *et al.* (1997) suggesting that Fe³⁺ in chlorite shows a strong preference for the M4 site where it substitutes for Al³⁺.

Several methods are available to measure Fe³⁺ in chlorite: chemical titration (Tschermak, 1891; Orcel, 1927; Hallimond *et al.*, 1939; Brindley, 1951), Mössbauer (Pal *et al.*, 1993; Aja & Dyar, 2000; Inoue *et al.*, 2009), X-ray photoelectron spectroscopy (XPS) (Inoue *et al.*, 2010) and micro-X-ray absorption near-edge spectroscopy (μ-XANES) (Wilke *et al.*,

2001; Vidal *et al.*, 2006; Rigault, 2010; Trincal *et al.*, 2015).

CHLORITE DATA FROM THE LITERATURE

A compilation of almost 200 published data of chlorite structural formulae containing Fe³⁺ is reported in Table 2. These chlorites come from a wide range of rocks from various localities. Some are of detrital origin; another experienced diagenetic to metamorphic conditions as indicated by the oscillatory zoned chlorites from the Pic-de-Port-Vieux outcrop (Fig. 1; Trincal *et al.*, 2015). In addition, 202 data from three localities were incorporated in this review (Inoue *et al.*, 2009). Most of the chlorites selected contain (in a.p.f.u, see Table 2): 2.5 to 3.5 Si⁴⁺, 2 to 4 Al³⁺ (Fig. 2), 1.5 to 5.5 R²⁺ (Mg²⁺, Mn²⁺, Fe²⁺) (Fig. 3) and <1 Fe³⁺ (Fig. 4). These analyses are distributed between classical end-members and apparently follow the classical DT and TK substitutions (Figs 2, 3) or the Al-Fe³⁺ substitution (Fig. 4).

In diagenetic to metamorphic rocks, the χFe^{3+} content of chlorite depends on the temperature and the oxygen fugacity that is controlled by the buffering assemblage, *i.e.* the bulk-rock composition (Lanari *et al.* 2014a). This control is supported by data from Inoue *et al.* (2009), which show relationships between sampling localities and chlorite structural formulae and oxidation state (Figs 2–4). It is therefore critical to look at the variability of χFe^{3+} recorded by chlorite in one rock sample, for a given bulk-rock composition. In such cases the observed variability of χFe^{3+} is caused by temperature variations,

Compilation of structural formulae for Fe³⁺-bearing chlorites, calculated on the basis of O = 14.

Mod	CaO + Na ₂ O +																	□	
	SiO ₂	TiO ₂	Al ₂ O ₃	Fe ₂ O ₃	FeO	MgO	MnO	K ₂ O	Others	Total	Si	Ti	^{IV} Al	^{VI} Al	Fe ³⁺	Fe ²⁺	Mg		Mn
	25.17	0.01	22.76		20.98	17.27	0.09	0.05	0.05	86.37	2.58		1.42	1.33	0.68	1.04	2.64		0.30
	26.11	0.01	22.93		18.43	19.64	0.09	0.04	0.05	87.29	2.63		1.37	1.35	0.38	1.13	2.95		0.18
	25.74	0.01	23.25		20.43	17.86	0.09	0.04	0.05	87.47	2.60		1.40	1.37	0.53	1.14	2.69		0.25
	25.98	0.01	23.65		18.39	19.09	0.09	0.04	0.05	87.30	2.62		1.38	1.42	0.32	1.19	2.87		0.18
	25.53	0.01	23.16		21.94	16.41	0.08	0.04	0.04	87.23	2.59		1.41	1.37	0.74	1.04	2.48		0.35
	25.76	0.02	22.91		21.92	16.36	0.10	0.05	0.05	87.17	2.62		1.38	1.37	0.70	1.08	2.48		0.35
	26.39	0.01	23.00		17.59	19.83	0.11	0.05	0.05	87.02	2.65		1.35	1.38	0.36	1.08	2.97		0.20
	25.75	0.02	23.17		20.47	17.44	0.11	0.05	0.05	87.04	2.62		1.38	1.39	0.53	1.15	2.64		0.27
	25.85	0.01	23.55		18.70	18.83	0.11	0.05	0.05	87.15	2.61		1.39	1.42	0.33	1.22	2.84		0.18
	25.55	0.02	23.05		21.48	16.58	0.11	0.05	0.05	86.87	2.60		1.40	1.37	0.72	1.02	2.52		0.35
	33.76		12.69		4.17	33.65			1.59	86.14	3.23		0.77	0.66	0.20	0.14	4.80		0.04
	33.73		13.54		3.35	33.92			1.40	87.00	3.19		0.81	0.70	0.17	0.16	4.79		0.03
	33.47		13.24		3.26	33.66			1.76	85.57	3.21		0.79	0.71	0.13	0.13	4.81		0.02
											2.67	0.01	1.33	1.24	0.47	1.51	2.75	0.01	0.19
	29.98		15.56	6.36	16.45	16.68	0.40	0.33		85.75	3.13		0.87	1.05	0.50	1.44	2.60	0.04	0.33
	28.08	0.07	18.34	3.80	13.82	19.08	1.12	0.31		84.60	2.93	0.01	1.07	1.18	0.30	1.21	2.97	0.10	0.19
	27.34	0.03	18.58	4.51	16.42	17.75	0.35	0.27		85.24	2.87	0.00	1.13	1.17	0.36	1.44	2.78	0.03	0.18
	29.98	0.04	19.05	3.91	11.26	20.30	0.60	0.29		84.45	3.03	0.00	0.97	1.30	0.30	0.95	3.06	0.05	0.30
	24.26	0.01	17.64	5.56	33.82	1.37	3.34	0.11		86.09	2.85	0.00	1.15	1.29	0.49	3.32	0.24	0.33	0.31
	27.17	0.05	18.77	3.76	19.37	13.71	3.21	0.24		86.29	2.89	0.00	1.11	1.25	0.30	1.73	2.18	0.29	0.21
	22.86	0.03	17.55	6.40	30.55	1.92	2.15	0.13		81.59	2.80	0.00	1.20	1.33	0.59	3.13	0.35	0.22	0.35
											2.63	0.01	1.36	1.41	0.13	2.51	1.84	0.01	0.08
											2.71		1.29	1.60	0.14	2.58	1.40	0.05	0.22
											2.92		1.08	1.62	0.10	1.82	2.11	0.02	0.31
											2.86		1.14	1.44	0.34	0.34	3.54		0.32
											2.78	0.01	1.21	1.10	0.79	0.79	2.94	0.03	0.34
											2.57	0.01	1.42	1.29	0.75	1.84	1.79	0.01	0.30
											2.93		1.07	1.53	0.18	0.39	3.57		0.32
											2.67	0.03	1.31	0.33	1.74	1.51	2.22	0.01	0.38
											2.76		1.24	1.37	0.49	2.14	1.62		0.31
											2.72		1.28	1.32	0.43	2.11	1.85		0.24
											2.71		1.29	1.31	0.14	2.58	1.84		0.07

(continued)

Al-free di-trioctahedral substitution in chlorite and a ferri-sudloite end-member

CaO + Na ₂ O +																	□
SiO ₂	TiO ₂	Al ₂ O ₃	Fe ₂ O ₃	FeO	MgO	MnO	K ₂ O	Others	Total	Si	Ti	^{IV} Al	^{VI} Al	Fe ³⁺	Fe ²⁺	Mg	
30.70	0.25	22.90	1.42		31.10	0.01	<0.20	0.01		2.69		1.31	1.31	0.14	2.60	1.82	0.07
33.60	0.11	13.70	5.95		32.90	0.13	<0.74	0.01		2.88		1.12	1.41	0.01	0.10	4.35	0.15
26.15		21.13	5.93	17.20	17.73				88.14	2.67		1.33	1.21	0.46	1.47	2.70	0.17
26.83		21.64	6.07	13.30	20.67				88.51	2.67		1.33	1.21	0.45	1.11	3.07	0.17
29.64	0.04	23.93		37.94	8.11	0.29	0.05	0.01		2.82		1.19	1.49	0.30	2.72	1.15	0.03
33.90	0.06	23.77		8.33	33.80	0.04	0.10	0.01		2.83		1.17	1.17	0.07	0.50	4.20	0.01
26.73	1.92	20.21		41.25	7.00	0.29	1.49	1.11		2.63	0.14	1.23	1.11	0.28	3.08	1.03	0.03
37.02		35.61	0.89		11.07	0.07	1.20			3.23		0.77	3.11	0.03	0.02	1.65	1.19
36.70		33.04	3.74		11.04		0.65			3.32		0.68	2.93	0.15	0.13	1.58	1.20
36.81		30.58	7.25		8.13	0.11	1.14			3.42		0.58	2.87	0.33	0.24	1.24	0.01
22.46	0.06	19.29	4.70	37.20	2.87	0.07			86.65	2.62	0.01	1.38	1.28	0.41	3.63	0.50	0.01
30.38	0.10	18.70	0.79	6.29	29.81	0.05			86.12	2.94	0.01	1.06	1.08	0.06	0.51	4.30	0.00
										2.68		1.32	1.18	0.18	1.95	2.68	0.02
27.69	0.02	22.26		12.26	24.56	0.17	0.01	0.08	87.05	2.73	0.00	1.27	1.32	0.37	0.64	3.61	0.01
26.74	0.06	21.77		11.72	24.36	0.17	0.01	0.17	85.00	2.70	0.01	1.29	1.31	0.37	0.62	3.67	0.02
27.42	0.05	22.11		12.28	24.72	0.15	0.01	0.16	85.67	2.72	0.00	1.28	1.30	0.38	0.64	3.65	0.01
31.68	0.05	17.13	1.34	1.62	34.76	0.08		0.05		2.99	0.00	1.01	0.89	0.10	0.13	4.88	0.01
										3.50		0.50	0.70	0.10	0.10	4.90	0.15
										3.00		1.00	0.90	0.10	0.10	4.90	0
										3.12		0.88	1.69	0.64	1.03	2.17	0.73
										3.01		0.99	0.49	0.29	0.50	4.20	0
										2.76		1.24	2.01	0.12	0.70	3.30	0.45
27.69	0.02	22.26		12.26	24.56	0.17	0.01	0.08	87.05	2.73	0.00	1.27	1.32	0.38	0.63	3.61	0.01
26.74	0.06	21.77		11.72	24.36	0.17	0.01	0.17	85.00	2.70	0.00	1.29	1.30	0.37	0.62	3.67	0.01
27.42	0.05	22.11		12.28	24.72	0.15	0.01	0.16	86.90	2.71	0.00	1.29	1.30	0.38	0.64	3.65	0.01
35.70		33.77	2.81	0.35	13.26	0.04		0.06	86.00	3.24		0.76	2.85	0.19	0.03	1.79	1.14
39.25	0.48	32.94	2.98		9.49		0.26	0.02		3.54	0.03	0.43	3.07	0.20		1.28	1.41
24.40		21.14		32.65	11.74	0.07				2.59		1.41	1.24	0.16	2.73	1.86	0.01
										2.85		1.15	0.97	0.18	0.28	4.55	0.00
										2.96		1.04	0.54	0.23	0.09	4.73	0
										2.83		1.17	4.32	0.07			1.61
										2.88		1.12	1.32	0.23	2.07	2.16	0.05
										2.68		1.32	1.06	0.14	1.44	3.06	0.07
										2.64		1.36	0.85	0.32	1.82	3.10	0

Sample	CaO + Na ₂ O +																		
	SiO ₂	TiO ₂	Al ₂ O ₃	Fe ₂ O ₃	FeO	MgO	MnO	K ₂ O	Others	Total	Si	Ti	^{IV} Al	^{VI} Al	Fe ³⁺	Fe ²⁺	Mg	Mn	□
											2.54		1.46	0.90	1.13	2.60	1.10		0.28
											2.49		1.51	1.51	0.61	2.53	0.71	0.25	0.31
											2.54	0.05	1.41	1.05	0.54	2.19	2.00	0.12	0.09
											2.55		1.45	1.37	0.33	1.50	2.65	0.03	0.13
											3.19		0.81	3.13	0.01	0.01	1.60		1.13
34.36		38.84	0.49			10.21		0.85		85.75	3.12		0.88	3.29	0.03		1.38		1.16
38.26		44.28	1.31	0.48					2.00		3.42		0.58	4.08	0.09	0.04			1.79
33.10	0.30	37.43	0.87			13.12		0.44			85.26	3.01	0.02	0.97	3.05	0.06		1.78	
29.45	0.20	20.14	4.08	18.69	14.93	0.36	1.59			89.34	2.97	0.02	1.02	1.36	0.31	1.58	2.24	0.03	0.27
30.84	0.12	14.15	2.59	18.42	20.62	0.10	0.84			87.68	3.15	0.01	0.84	0.87	0.20	1.58	3.14	0.01	0.09
25.80		18.30	4.10	21.30	20.30					89.80	2.64		1.36	0.85	0.32	1.82	3.10		0
27.90		25.20		10.60	24.20					87.90	2.69		1.31	1.55		0.85	3.48		0.12
30.50		17.30	2.90	6.10	32.80					89.60	2.87		1.13	0.79	0.21	0.48	4.60		0
33.90		14.60	2.70	2.10	34.60					87.90	3.16		0.84	0.77	0.19	0.16	4.81		0.06
											3.14		0.86	4.10	0.04	0.01	0.08		1.64
27.27		15.21	4.35	2.78	10.13	0.06	0.28		29.87	89.98	2.99		1.01	0.95	0.36	0.25	1.65	0.01	0.15
27.27		15.21	4.35	2.78	10.13	0.06	0.38		29.87		2.98		1.02	0.95	0.36	0.25	1.65	0.01	0.14
32.72	0.03	20.28	1.00		24.47	0.06	2.22			80.78	3.24	0.00	0.76	1.61	0.07		3.61	0.01	0.44
31.17		37.49	1.04	1.95	14.25		0.26				2.85		1.15	2.90	0.07	0.15	1.95		0.91
											3.45		0.55	3.96	0.02	0.02	0.04	0.01	1.72
											2.66		1.34	1.74	0.02	3.68	0.42		0.21
											3.43		0.57	2.70	0.09	0.04	1.97		1.11
27.31	0.22	19.69	3.23	18.58	19.01	0.27	0.80			89.11	2.78	0.02	1.20	1.16	0.25	1.58	2.88	0.02	0.11
35.63		34.87	5.01	0.43	8.63	0.05	1.83			86.45	3.26		0.74	3.02	0.34	0.04	1.17		1.31
39.01	0.47	32.15	0.90	0.10	10.14		2.16				3.56	0.03	0.41	3.05	0.06	0.01	1.37		1.25
30.66		30.55	0.47		10.13		1.79				3.24		0.76	3.04	0.04		1.59		1.00
24.45	0.21	21.99	20.80	28.04	11.90	0.54	0.25			89.46	2.59	0.02	1.39	1.36	0.17	2.19	1.88	0.05	0.06
											2.70		1.30	4.30	0.03	0.02			1.52
											2.80	0.09	1.11	3.31	0.38		0.76		1.18
28.32	0.09	19.03	1.19	14.85	23.72	0.09	0.63			87.92	2.83	0.01	1.17	1.06	0.09	1.24	3.53	0.01	0
21.34	0.21	20.73	5.20	40.17	0.99	0.89	0.04			89.57	2.46	0.02	1.52	1.30	0.45	3.87	0.17	0.09	0.12
36.43		12.24	0.94	6.87	30.94	0.11	0.33		0.63	88.49	3.43		0.57	0.79	0.07	0.54	4.35	0.01	0.15
33.14	0.08	10.04	4.00	5.66	32.64		0.80				86.36	3.25	0.01	0.75	0.41	0.29	0.46	4.77	
29.73		17.95	0.68	10.05	21.84	8.24				88.49	2.98		1.02	1.09	0.05	0.84	3.26	0.70	0.06
22.81		20.35	4.84	34.87	6.11	0.56	0.10				2.54		1.46	1.20	0.40	3.24	1.01	0.05	0.07

23.38	19.32	2.51	38.38	5.39	0.58			89.56	2.63	1.37	1.20	0.21	3.61	0.90	0.06	0.02
26.04	19.96	1.85	21.34	18.56	0.47			88.22	2.69	1.31	1.13	0.14	1.85	2.86	0.04	0
29.71	19.75	0.29	6.59	30.79	0.30			87.43	2.84	1.16	1.07	0.02	0.53	4.39	0.02	0
32.27	14.99	1.56	6.89	31.75	0.09			87.55	3.10	0.90	0.79	0.11	0.55	4.54	0.01	0.00
27.11	0.35	17.42	2.91	30.98	9.75	0.21		88.73	2.93	0.03	1.07	1.15	0.24	2.80	1.57	0.16
29.94	0.16	13.77	10.46	20.96	10.71	0.18	1.34	87.52	3.19	0.01	0.81	0.92	0.84	1.87	1.70	0.02
28.73	0.41	19.16	1.97	10.99	26.37	0.15	0.10	87.88	2.81	0.03	1.19	1.03	0.15	0.90	3.85	0.01
22.47		21.82	0.22	37.24	2.57			84.32	2.65		1.35	1.68	0.02	3.67	0.45	0.18
23.48		22.45	43.37	0.45	2.60			92.35	2.35		1.65	1.00	3.27	0.04	0.39	1.31
33.40		47.47		0.71	0.20	0.54		3.12	85.44	3.07	0.93	4.22		0.05	0.03	1.64
29.84	0.15	20.20	0.80	15.00	21.83	0.13	0.80	0.03	88.78	2.93	0.01	1.07	1.27	0.06	1.23	3.19
22.03	3.63	22.91	0.46	36.68	1.91	0.04	0.18	0.26	88.77	2.47	0.31	1.53	1.50	0.04	3.44	0.32
									3.10			0.90	1.30	0.60	3.35	0.75
									2.48	0.30	1.22	1.82	0.04	3.46	0.32	0.32
21.40	0.89	25.40	0.25	37.60	2.04	0.05	0.13		87.98	2.42	0.08	1.58	1.80	0.02	3.55	0.34
									2.48	0.30	1.22	1.82	0.04	3.46	0.32	0.32
26.40		18.23	5.70	25.87	11.35	0.04	0.76		88.35	2.83		1.17	1.13	0.46	2.32	1.81
27.64	0.22	22.48	0.06	12.06	24.32	0.02	0.23	0.06	87.09	2.73	0.02	1.27	1.34	0.00	0.99	3.57
31.87	0.17	14.51	1.86	3.57	32.76		0.06	1.38	86.18	3.07	0.01	0.93	0.72	0.14	0.29	4.71
20.82		17.64	8.70	37.96	4.15				89.27	2.42		1.58	0.83	0.76	3.69	0.72
26.65		16.14	6.69	34.43	4.47				88.38	2.99		1.01	1.12	0.56	3.23	0.75
22.47		23.57	4.01	29.27	9.81				89.13	2.42		1.58	1.40	0.32	2.63	1.57
26.45		20.88	2.82	21.06	16.84	0.44	0.47		88.96	2.71		1.29	1.24	0.22	1.81	2.57
									3.09			0.91	3.92	0.05	0.21	1.44
									3.06			0.94	3.88	0.08	0.23	1.40
									3.10			0.90	4.16	0.07	0.11	1.67
									2.80			1.20	0.86	0.60	3.56	0.78
24.35	0.04	20.21	2.13	36.27	5.57	0.48	0.10		89.15	2.70	0.00	1.30	1.35	0.18	3.37	0.92
24.90		15.60	7.20	35.00	4.60	0.40			87.70	2.86		1.14	0.97	0.62	3.36	0.79
25.07	0.12	19.78	3.50	35.80	1.11	0.50	2.15		88.03	2.85	0.01	1.15	1.50	0.30	3.40	1.19
25.09	0.05	18.94	9.60	25.66	9.80	0.02	0.49		89.65	2.68	0.00	1.32	1.06	0.77	2.29	1.56
22.18	0.04	20.04	7.35	35.23	3.79	0.02	0.49		90.60	2.51	0.00	1.49	1.18	0.63	3.33	0.64
26.69	0.30	19.57	3.49	21.80	16.23	0.30	0.17	0.13	88.68	2.76	0.02	1.24	1.15	0.27	1.89	2.50
27.56		24.47	3.85	10.51	20.86	1.80			89.05	2.68		1.32	1.48	0.28	0.85	3.02
25.62	0.88	21.19	3.88	21.55	15.28	0.35	0.16		88.91	2.64	0.07	1.36	1.22	0.30	1.86	2.35
23.32	0.03	17.45	4.09	38.90	4.54	0.01	0.24		88.58	2.68	0.00	1.32	1.05	0.35	3.74	0.78
23.43		18.21	4.49	36.30	5.41				87.84	2.68		1.32	1.13	0.39	3.47	0.92
24.60		19.10	3.12	29.04	13.10				88.96	2.64		1.36	1.06	0.25	2.61	2.10

(continued)

Sample	Chemical composition (wt %)														Fe speciation (at %)				Total Fe	Fe ²⁺	Fe ³⁺	Fe ⁴⁺	Fe ⁵⁺	Fe ⁶⁺	Fe ⁷⁺	Fe ⁸⁺	Fe ⁹⁺	Fe ¹⁰⁺	Fe ¹¹⁺	Fe ¹²⁺	Fe ¹³⁺	Fe ¹⁴⁺	Fe ¹⁵⁺	Fe ¹⁶⁺	Fe ¹⁷⁺	Fe ¹⁸⁺	Fe ¹⁹⁺	Fe ²⁰⁺	Fe ²¹⁺	Fe ²²⁺	Fe ²³⁺	Fe ²⁴⁺	Fe ²⁵⁺	Fe ²⁶⁺	Fe ²⁷⁺	Fe ²⁸⁺	Fe ²⁹⁺	Fe ³⁰⁺	Fe ³¹⁺	Fe ³²⁺	Fe ³³⁺	Fe ³⁴⁺	Fe ³⁵⁺	Fe ³⁶⁺	Fe ³⁷⁺	Fe ³⁸⁺	Fe ³⁹⁺	Fe ⁴⁰⁺	Fe ⁴¹⁺	Fe ⁴²⁺	Fe ⁴³⁺	Fe ⁴⁴⁺	Fe ⁴⁵⁺	Fe ⁴⁶⁺	Fe ⁴⁷⁺	Fe ⁴⁸⁺	Fe ⁴⁹⁺	Fe ⁵⁰⁺	Fe ⁵¹⁺	Fe ⁵²⁺	Fe ⁵³⁺	Fe ⁵⁴⁺	Fe ⁵⁵⁺	Fe ⁵⁶⁺	Fe ⁵⁷⁺	Fe ⁵⁸⁺	Fe ⁵⁹⁺	Fe ⁶⁰⁺	Fe ⁶¹⁺	Fe ⁶²⁺	Fe ⁶³⁺	Fe ⁶⁴⁺	Fe ⁶⁵⁺	Fe ⁶⁶⁺	Fe ⁶⁷⁺	Fe ⁶⁸⁺	Fe ⁶⁹⁺	Fe ⁷⁰⁺	Fe ⁷¹⁺	Fe ⁷²⁺	Fe ⁷³⁺	Fe ⁷⁴⁺	Fe ⁷⁵⁺	Fe ⁷⁶⁺	Fe ⁷⁷⁺	Fe ⁷⁸⁺	Fe ⁷⁹⁺	Fe ⁸⁰⁺	Fe ⁸¹⁺	Fe ⁸²⁺	Fe ⁸³⁺	Fe ⁸⁴⁺	Fe ⁸⁵⁺	Fe ⁸⁶⁺	Fe ⁸⁷⁺	Fe ⁸⁸⁺	Fe ⁸⁹⁺	Fe ⁹⁰⁺	Fe ⁹¹⁺	Fe ⁹²⁺	Fe ⁹³⁺	Fe ⁹⁴⁺	Fe ⁹⁵⁺	Fe ⁹⁶⁺	Fe ⁹⁷⁺	Fe ⁹⁸⁺	Fe ⁹⁹⁺	Fe ¹⁰⁰⁺	Fe ¹⁰¹⁺	Fe ¹⁰²⁺	Fe ¹⁰³⁺	Fe ¹⁰⁴⁺	Fe ¹⁰⁵⁺	Fe ¹⁰⁶⁺	Fe ¹⁰⁷⁺	Fe ¹⁰⁸⁺	Fe ¹⁰⁹⁺	Fe ¹¹⁰⁺	Fe ¹¹¹⁺	Fe ¹¹²⁺	Fe ¹¹³⁺	Fe ¹¹⁴⁺	Fe ¹¹⁵⁺	Fe ¹¹⁶⁺	Fe ¹¹⁷⁺	Fe ¹¹⁸⁺	Fe ¹¹⁹⁺	Fe ¹²⁰⁺	Fe ¹²¹⁺	Fe ¹²²⁺	Fe ¹²³⁺	Fe ¹²⁴⁺	Fe ¹²⁵⁺	Fe ¹²⁶⁺	Fe ¹²⁷⁺	Fe ¹²⁸⁺	Fe ¹²⁹⁺	Fe ¹³⁰⁺	Fe ¹³¹⁺	Fe ¹³²⁺	Fe ¹³³⁺	Fe ¹³⁴⁺	Fe ¹³⁵⁺	Fe ¹³⁶⁺	Fe ¹³⁷⁺	Fe ¹³⁸⁺	Fe ¹³⁹⁺	Fe ¹⁴⁰⁺	Fe ¹⁴¹⁺	Fe ¹⁴²⁺	Fe ¹⁴³⁺	Fe ¹⁴⁴⁺	Fe ¹⁴⁵⁺	Fe ¹⁴⁶⁺	Fe ¹⁴⁷⁺	Fe ¹⁴⁸⁺	Fe ¹⁴⁹⁺	Fe ¹⁵⁰⁺	Fe ¹⁵¹⁺	Fe ¹⁵²⁺	Fe ¹⁵³⁺	Fe ¹⁵⁴⁺	Fe ¹⁵⁵⁺	Fe ¹⁵⁶⁺	Fe ¹⁵⁷⁺	Fe ¹⁵⁸⁺	Fe ¹⁵⁹⁺	Fe ¹⁶⁰⁺	Fe ¹⁶¹⁺	Fe ¹⁶²⁺	Fe ¹⁶³⁺	Fe ¹⁶⁴⁺	Fe ¹⁶⁵⁺	Fe ¹⁶⁶⁺	Fe ¹⁶⁷⁺	Fe ¹⁶⁸⁺	Fe ¹⁶⁹⁺	Fe ¹⁷⁰⁺	Fe ¹⁷¹⁺	Fe ¹⁷²⁺	Fe ¹⁷³⁺	Fe ¹⁷⁴⁺	Fe ¹⁷⁵⁺	Fe ¹⁷⁶⁺	Fe ¹⁷⁷⁺	Fe ¹⁷⁸⁺	Fe ¹⁷⁹⁺	Fe ¹⁸⁰⁺	Fe ¹⁸¹⁺	Fe ¹⁸²⁺	Fe ¹⁸³⁺	Fe ¹⁸⁴⁺	Fe ¹⁸⁵⁺	Fe ¹⁸⁶⁺	Fe ¹⁸⁷⁺	Fe ¹⁸⁸⁺	Fe ¹⁸⁹⁺	Fe ¹⁹⁰⁺	Fe ¹⁹¹⁺	Fe ¹⁹²⁺	Fe ¹⁹³⁺	Fe ¹⁹⁴⁺	Fe ¹⁹⁵⁺	Fe ¹⁹⁶⁺	Fe ¹⁹⁷⁺	Fe ¹⁹⁸⁺	Fe ¹⁹⁹⁺	Fe ²⁰⁰⁺	Fe ²⁰¹⁺	Fe ²⁰²⁺	Fe ²⁰³⁺	Fe ²⁰⁴⁺	Fe ²⁰⁵⁺	Fe ²⁰⁶⁺	Fe ²⁰⁷⁺	Fe ²⁰⁸⁺	Fe ²⁰⁹⁺	Fe ²¹⁰⁺	Fe ²¹¹⁺	Fe ²¹²⁺	Fe ²¹³⁺	Fe ²¹⁴⁺	Fe ²¹⁵⁺	Fe ²¹⁶⁺	Fe ²¹⁷⁺	Fe ²¹⁸⁺	Fe ²¹⁹⁺	Fe ²²⁰⁺	Fe ²²¹⁺	Fe ²²²⁺	Fe ²²³⁺	Fe ²²⁴⁺	Fe ²²⁵⁺	Fe ²²⁶⁺	Fe ²²⁷⁺	Fe ²²⁸⁺	Fe ²²⁹⁺	Fe ²³⁰⁺	Fe ²³¹⁺	Fe ²³²⁺	Fe ²³³⁺	Fe ²³⁴⁺	Fe ²³⁵⁺	Fe ²³⁶⁺	Fe ²³⁷⁺	Fe ²³⁸⁺	Fe ²³⁹⁺	Fe ²⁴⁰⁺	Fe ²⁴¹⁺	Fe ²⁴²⁺	Fe ²⁴³⁺	Fe ²⁴⁴⁺	Fe ²⁴⁵⁺	Fe ²⁴⁶⁺	Fe ²⁴⁷⁺	Fe ²⁴⁸⁺	Fe ²⁴⁹⁺	Fe ²⁵⁰⁺	Fe ²⁵¹⁺	Fe ²⁵²⁺	Fe ²⁵³⁺	Fe ²⁵⁴⁺	Fe ²⁵⁵⁺	Fe ²⁵⁶⁺	Fe ²⁵⁷⁺	Fe ²⁵⁸⁺	Fe ²⁵⁹⁺	Fe ²⁶⁰⁺	Fe ²⁶¹⁺	Fe ²⁶²⁺	Fe ²⁶³⁺	Fe ²⁶⁴⁺	Fe ²⁶⁵⁺	Fe ²⁶⁶⁺	Fe ²⁶⁷⁺	Fe ²⁶⁸⁺	Fe ²⁶⁹⁺	Fe ²⁷⁰⁺	Fe ²⁷¹⁺	Fe ²⁷²⁺	Fe ²⁷³⁺	Fe ²⁷⁴⁺	Fe ²⁷⁵⁺	Fe ²⁷⁶⁺	Fe ²⁷⁷⁺	Fe ²⁷⁸⁺	Fe ²⁷⁹⁺	Fe ²⁸⁰⁺	Fe ²⁸¹⁺	Fe ²⁸²⁺	Fe ²⁸³⁺	Fe ²⁸⁴⁺	Fe ²⁸⁵⁺	Fe ²⁸⁶⁺	Fe ²⁸⁷⁺	Fe ²⁸⁸⁺	Fe ²⁸⁹⁺	Fe ²⁹⁰⁺	Fe ²⁹¹⁺	Fe ²⁹²⁺	Fe ²⁹³⁺	Fe ²⁹⁴⁺	Fe ²⁹⁵⁺	Fe ²⁹⁶⁺	Fe ²⁹⁷⁺	Fe ²⁹⁸⁺	Fe ²⁹⁹⁺	Fe ³⁰⁰⁺	Fe ³⁰¹⁺	Fe ³⁰²⁺	Fe ³⁰³⁺	Fe ³⁰⁴⁺	Fe ³⁰⁵⁺	Fe ³⁰⁶⁺	Fe ³⁰⁷⁺	Fe ³⁰⁸⁺	Fe ³⁰⁹⁺	Fe ³¹⁰⁺	Fe ³¹¹⁺	Fe ³¹²⁺	Fe ³¹³⁺	Fe ³¹⁴⁺	Fe ³¹⁵⁺	Fe ³¹⁶⁺	Fe ³¹⁷⁺	Fe ³¹⁸⁺	Fe ³¹⁹⁺	Fe ³²⁰⁺	Fe ³²¹⁺	Fe ³²²⁺	Fe ³²³⁺	Fe ³²⁴⁺	Fe ³²⁵⁺	Fe ³²⁶⁺	Fe ³²⁷⁺	Fe ³²⁸⁺	Fe ³²⁹⁺	Fe ³³⁰⁺	Fe ³³¹⁺	Fe ³³²⁺	Fe ³³³⁺	Fe ³³⁴⁺	Fe ³³⁵⁺	Fe ³³⁶⁺	Fe ³³⁷⁺	Fe ³³⁸⁺	Fe ³³⁹⁺	Fe ³⁴⁰⁺	Fe ³⁴¹⁺	Fe ³⁴²⁺	Fe ³⁴³⁺	Fe ³⁴⁴⁺	Fe ³⁴⁵⁺	Fe ³⁴⁶⁺	Fe ³⁴⁷⁺	Fe ³⁴⁸⁺	Fe ³⁴⁹⁺	Fe ³⁵⁰⁺	Fe ³⁵¹⁺	Fe ³⁵²⁺	Fe ³⁵³⁺	Fe ³⁵⁴⁺	Fe ³⁵⁵⁺	Fe ³⁵⁶⁺	Fe ³⁵⁷⁺	Fe ³⁵⁸⁺	Fe ³⁵⁹⁺	Fe ³⁶⁰⁺	Fe ³⁶¹⁺	Fe ³⁶²⁺	Fe ³⁶³⁺	Fe ³⁶⁴⁺	Fe ³⁶⁵⁺	Fe ³⁶⁶⁺	Fe ³⁶⁷⁺	Fe ³⁶⁸⁺	Fe ³⁶⁹⁺	Fe ³⁷⁰⁺	Fe ³⁷¹⁺	Fe ³⁷²⁺	Fe ³⁷³⁺	Fe ³⁷⁴⁺	Fe ³⁷⁵⁺	Fe ³⁷⁶⁺	Fe ³⁷⁷⁺	Fe ³⁷⁸⁺	Fe ³⁷⁹⁺	Fe ³⁸⁰⁺	Fe ³⁸¹⁺	Fe ³⁸²⁺	Fe ³⁸³⁺	Fe ³⁸⁴⁺	Fe ³⁸⁵⁺	Fe ³⁸⁶⁺	Fe ³⁸⁷⁺	Fe ³⁸⁸⁺	Fe ³⁸⁹⁺	Fe ³⁹⁰⁺	Fe ³⁹¹⁺	Fe ³⁹²⁺	Fe ³⁹³⁺	Fe ³⁹⁴⁺	Fe ³⁹⁵⁺	Fe ³⁹⁶⁺	Fe ³⁹⁷⁺	Fe ³⁹⁸⁺	Fe ³⁹⁹⁺	Fe ⁴⁰⁰⁺	Fe ⁴⁰¹⁺	Fe ⁴⁰²⁺	Fe ⁴⁰³⁺	Fe ⁴⁰⁴⁺	Fe ⁴⁰⁵⁺	Fe ⁴⁰⁶⁺	Fe ⁴⁰⁷⁺	Fe ⁴⁰⁸⁺	Fe ⁴⁰⁹⁺	Fe ⁴¹⁰⁺	Fe ⁴¹¹⁺	Fe ⁴¹²⁺	Fe ⁴¹³⁺	Fe ⁴¹⁴⁺	Fe ⁴¹⁵⁺	Fe ⁴¹⁶⁺	Fe ⁴¹⁷⁺	Fe ⁴¹⁸⁺	Fe ⁴¹⁹⁺	Fe ⁴²⁰⁺	Fe ⁴²¹⁺	Fe ⁴²²⁺	Fe ⁴²³⁺	Fe ⁴²⁴⁺	Fe ⁴²⁵⁺	Fe ⁴²⁶⁺	Fe ⁴²⁷⁺	Fe ⁴²⁸⁺	Fe ⁴²⁹⁺	Fe ⁴³⁰⁺	Fe ⁴³¹⁺	Fe ⁴³²⁺	Fe ⁴³³⁺	Fe ⁴³⁴⁺	Fe ⁴³⁵⁺	Fe ⁴³⁶⁺	Fe ⁴³⁷⁺	Fe ⁴³⁸⁺	Fe ⁴³⁹⁺	Fe ⁴⁴⁰⁺	Fe ⁴⁴¹⁺	Fe ⁴⁴²⁺	Fe ⁴⁴³⁺	Fe ⁴⁴⁴⁺	Fe ⁴⁴⁵⁺	Fe ⁴⁴⁶⁺	Fe ⁴⁴⁷⁺	Fe ⁴⁴⁸⁺	Fe ⁴⁴⁹⁺	Fe ⁴⁵⁰⁺	Fe ⁴⁵¹⁺	Fe ⁴⁵²⁺	Fe ⁴⁵³⁺	Fe ⁴⁵⁴⁺	Fe ⁴⁵⁵⁺	Fe ⁴⁵⁶⁺	Fe ⁴⁵⁷⁺	Fe ⁴⁵⁸⁺	Fe ⁴⁵⁹⁺	Fe ⁴⁶⁰⁺	Fe ⁴⁶¹⁺	Fe ⁴⁶²⁺	Fe ⁴⁶³⁺	Fe ⁴⁶⁴⁺	Fe ⁴⁶⁵⁺	Fe ⁴⁶⁶⁺	Fe ⁴⁶⁷⁺	Fe ⁴⁶⁸⁺	Fe ⁴⁶⁹⁺	Fe ⁴⁷⁰⁺	Fe ⁴⁷¹⁺	Fe ⁴⁷²⁺	Fe ⁴⁷³⁺	Fe ⁴⁷⁴⁺	Fe ⁴⁷⁵⁺	Fe ⁴⁷⁶⁺	Fe ⁴⁷⁷⁺	Fe ⁴⁷⁸⁺	Fe ⁴⁷⁹⁺	Fe ⁴⁸⁰⁺	Fe ⁴⁸¹⁺	Fe ⁴⁸²⁺	Fe ⁴⁸³⁺	Fe ⁴⁸⁴⁺	Fe ⁴⁸⁵⁺	Fe ⁴⁸⁶⁺	Fe ⁴⁸⁷⁺	Fe ⁴⁸⁸⁺	Fe ⁴⁸⁹⁺	Fe ⁴⁹⁰⁺	Fe ⁴⁹¹⁺	Fe ⁴⁹²⁺	Fe ⁴⁹³⁺	Fe ⁴⁹⁴⁺	Fe ⁴⁹⁵⁺	Fe ⁴⁹⁶⁺	Fe ⁴⁹⁷⁺	Fe ⁴⁹⁸⁺	Fe ⁴⁹⁹⁺	Fe ⁵⁰⁰⁺	Fe ⁵⁰¹⁺	Fe ⁵⁰²⁺	Fe ⁵⁰³⁺	Fe ⁵⁰⁴⁺	Fe ⁵⁰⁵⁺	Fe ⁵⁰⁶⁺	Fe ⁵⁰⁷⁺	Fe ⁵⁰⁸⁺
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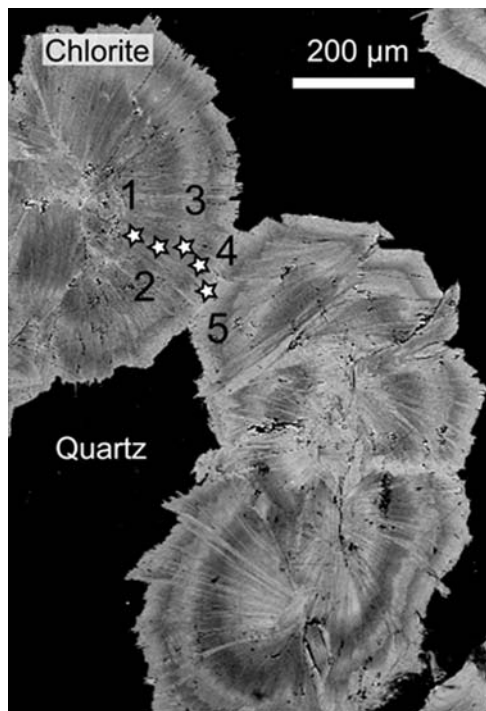


FIG. 1. Backscattered-electron image of an oscillatory-zoned chlorite showing Al-free di-trioctahedral substitution, from the Pic-de-Port-Vieux, Pyrenees, Spain. The numbers refer to the structural formulae reported in Table 2.

rather than by changes in the buffering assemblage (Lanari *et al.* 2014a; Trincal *et al.*, 2015). Most of the spot analyses reported in the literature correspond to different geological environments (Table 2); it is thus very difficult to use them to evaluate the link between the Fe^{3+} behaviour and the evolution of a parameter such as pressure, temperature, f_{O_2} , etc.

The choice of analytical method to estimate the $X\text{Fe}^{3+}$ in chlorite is essential. Chlorite Fe^{3+} measurements by Mössbauer spectroscopy or titration are relatively common but provide few constraints due to poor spatial resolution. Indeed, recent investigations using high-resolution mapping (*e.g.* de Andrade *et al.*, 2006; Muñoz *et al.*, 2006) have demonstrated that chlorite is frequently zoned, recording strong compositional variability even at the crystal scale (*e.g.* Lanari *et al.*, 2014b). Compositional zoning reflects time series of equilibrium conditions experienced by the rock. Thus *in situ* micrometric analyses are required to ensure a precise determination of the $\text{Fe}^{3+}/\text{Fe}_{\text{Tot}}$ of the successive growth zones, which is impossible using Mössbauer spectroscopy. μ -XANES $X\text{Fe}^{3+}$ spot analyses or

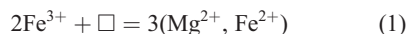
mapping allow us to elucidate the heterogeneity of the Fe chemistry of chlorite which is critical to see the relations between formation temperature and crystal geometry (Vidal *et al.*, 2006; Trincal *et al.*, 2015).

μ XANES DATA FROM ZONED CHLORITES

Accurate *in situ* μ XANES and electron microprobe analyses performed on zoned chlorite from the Sambagawa metamorphic belt in Japan (Vidal *et al.*, 2006) and on oscillatory-zoned chlorite from hydrothermal veins in the Pic-de-Port-Vieux, Pyrenees, Spain (fig. 1 from Trincal *et al.*, 2015) show variations in the composition of the Fe^{3+} , Mg and Fe^{2+} cation proportions, while that of Si and Al remain constant (Table 2, Fig. 2). This trend is observed independently in both samples, which have different bulk-rock compositions and experienced different metamorphic conditions. Indeed, chlorite from Japan comes from a blueschist-facies metapelite made of K-white mica, garnet, albite, quartz and graphite (Vidal *et al.*, 2006), while the chlorite from the Pyrenees crystallizes in a vein of a greenschist-facies metapelite without garnet or graphite (Trincal *et al.*, 2015). The absence of graphite is a good indicator of more oxidizing conditions. In both cases, $X\text{Fe}^{3+}$ increases as temperature decreases, as suggested by numerous authors (*e.g.* Lanari *et al.*, 2014a; Vidal *et al.*, 2016). Furthermore, in both cases, there is no significant change in the Al content of the zoned chlorite. This result excludes the hypothesis of DT, TK or Al- Fe^{3+} substitutions being at the origin of the $X\text{Fe}^{3+}$ variations.

AL-FREE DI-TRIOCTAHEDRAL SUBSTITUTION

Based on μ XANES analyses performed on zoned chlorite by Vidal *et al.* (2006) and Trincal *et al.* (2015), and in agreement with the literature data (Table 2), we demonstrate that a di-trioctahedral substitution implying Fe^{3+} can be used to explain the variations of $X\text{Fe}^{3+}$ in natural chlorite:



As this substitution does not explicitly involve any Al cations, it is named Al-free di-trioctahedral substitution (AFDT); it must not be confused with the Al-free chlorite end-member.

This substitution is similar to another couple suggested by Billault *et al.* (2002) and based on electron microprobe

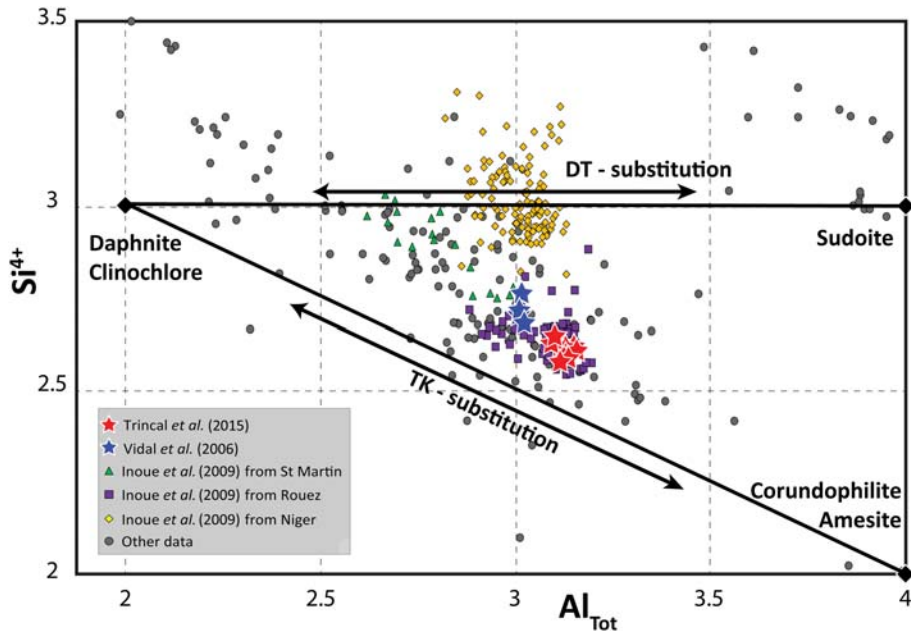


FIG. 2. Compilation of the Fe^{3+} -bearing chlorite compositions. The Si-Al plot is used to highlight the di-trioctahedral (DT) and Tschermak (TK) substitutions. End-members and structural formulae are reported in Tables 1 and 2.

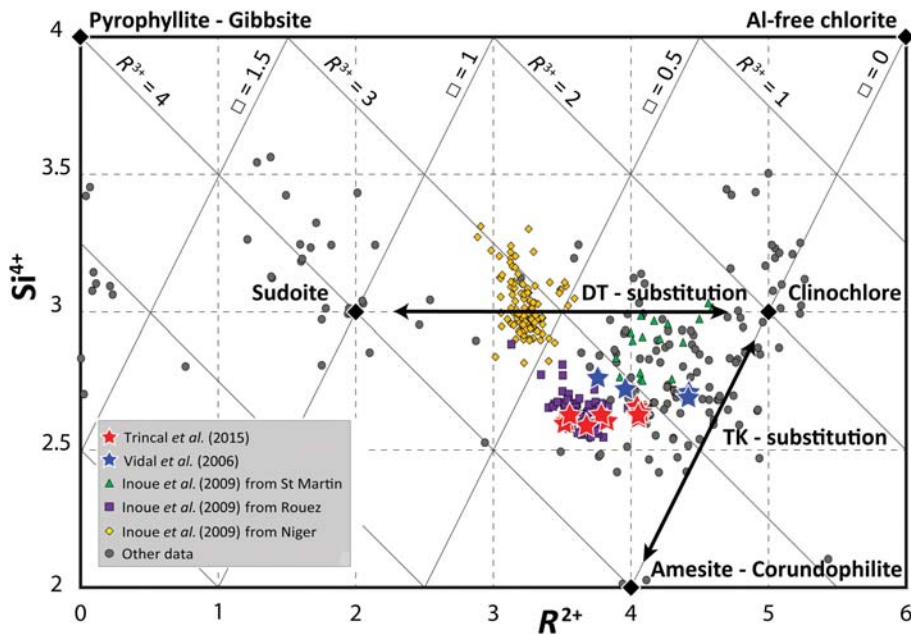


FIG. 3. Compilation of Fe^{3+} -bearing chlorite compositions. The Si^{4+} - R^{2+} plot by Wiewióra & Weiss (1990) is used to highlight the di-trioctahedral (DT) and the Al- Fe^{3+} substitutions.

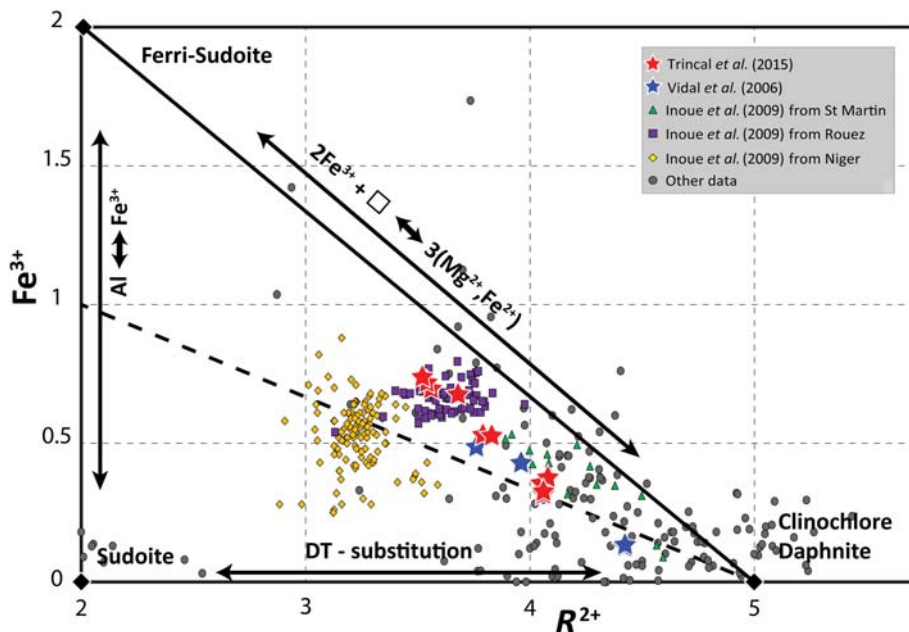


FIG. 4. Compilation of the Fe^{3+} -bearing chlorite compositions. The Fe^{3+} – R^{2+} plot highlights the di-trioctahedral (DT), Al- Fe^{3+} and AFDT substitutions.

and Mössbauer spectroscopic analyses on sudoite:

$$\text{Fe}^{3+} = \text{Al}_{(\text{VI})} \text{ and } \text{Fe}^{2+} + \text{Fe}^{3+} + \square = 3(\text{Mg}^{2+}) \quad (2)$$

However, chlorite analyses from their study display both Al and XMg variations, which is not the case for the samples in the present study.

Chlorite compositions from Vidal *et al.* (2006) and Trincal *et al.* (2015) are plotted in an Fe^{3+} vs. R^{2+} diagram (Fig. 4). In this diagram, data from zoned chlorite are distributed along a line parallel to the AFDT substitution vector. The AFDT substitution allows us to define a theoretical Fe^{3+} -bearing chlorite end-member; namely ferri-sudoite $((\text{Si}_3\text{Al})[(\text{Fe}^{2+}, \text{Mg})_2\text{Fe}_2^{3+}\square\text{Al}]\text{O}_{10}(\text{OH})_8)$ with two atoms of Fe^{3+} (Table 1). Ferri-sudoite can be separated into ferri-sudoite-Fe $((\text{Si}_3\text{Al})[\text{Fe}_2^{2+}\text{Fe}_2^{3+}\square\text{Al}]\text{O}_{10}(\text{OH})_8)$ and ferri-sudoite-Mg $((\text{Si}_3\text{Al})[\text{Mg}_2\text{Fe}_2^{3+}\square\text{Al}]\text{O}_{10}(\text{OH})_8)$. As Fe^{3+} replaces Mg and Fe^{2+} , it seems convenient to allocate up two atoms of Fe^{3+} to the M2-M3 sites (total multiplicity of 4) leaving Al in the M4 (Table 1).

According to several authors, the Fe^{3+} cation seems limited to 1 a.p.f.u and is generally allocated to the M4 site following the Al- Fe^{3+} substitution. This last assertion is supported by Z-contrast images of high-angle annular dark-field imaging-scanning

transmission electron microscopy (HAADF-STEM) and other studies (e.g. Smyth *et al.*, 1997; Inoue & Kogure, 2016; Vidal *et al.*, 2016). Among data collected in the literature, only six analyses show Fe^{3+} contents of >1 a.p.f.u (Orcel, 1927; Brindley & Youell, 1953; Malysheva *et al.*, 1977; Goodman & Bain, 1979; Kodama *et al.*, 1982; Singer *et al.*, 2009 in Table 2). However, these analyses would actually match other phyllosilicates, suggesting contamination issues. Although AFDT substitution suggests that it is possible, no sudoite with >1 Fe^{3+} atom has been yet discovered; it offers interesting research opportunities.

CONCLUSION

The compilation of Fe^{3+} -bearing chlorite analyses in this study, especially those acquired by μXANES spectroscopy in zoned crystals, suggests the existence of a new chlorite substitution, the AFDT substitution that is characterized by $2\text{Fe}^{3+} + \square = 3(\text{Mg}^{2+}, \text{Fe}^{2+})$ and implies a new chlorite end-member: ferri-sudoite $((\text{Si}_3\text{Al})[(\text{Fe}^{2+}, \text{Mg})_2\text{Fe}_2^{3+}\square\text{Al}]\text{O}_{10}(\text{OH})_8)$ with 2 atoms of Fe^{3+} . These results are not in agreement with previous studies which showed Al- Fe^{3+} substitution implying only 1 Fe^{3+} . The absence of constraints on the Fe^{3+} crystallographic position together with the Fe^{3+} ionic properties in chlorite require further studies to confirm

the existence of the AFDT substitution and of the ferri-sudoite end-member, *e.g.* with accurate *in situ* Fe³⁺ measurements in synthetic or diagenetic chlorite formed in different oxidizing environments. This study opens the door for further research developments that would have strong implications, *e.g.* in chlorite geothermometry.

ACKNOWLEDGEMENTS

The authors thank Daniel Beaufort, Olivier Vidal and Martin Engi for fruitful discussions regarding Fe³⁺ in chlorite; Atsuyuki Inoue, Fernando Nieto-Garcia and another referee for the valuable advice about the manuscript; and Flavien Choulet for corrections to the language.

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